INVESTIGACIÓN

A simple model of the diffusion phenomena taking place during the debittering process of green table olives

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RESUMEN

Un modelo simple del fenómeno de difusión que tiene lugar durante el proceso de desamarizado de aceitunas verdes de mesa.

Se ha cuantificado el cambio en la concentración de sodio y calcio en la pulpa y en la lejía durante el tratamiento alcalino de aceitunas. La concentración promedio de sodio en la pulpa aumentó de 0,0045 a 0,395 meq Na/g mientras que la de calcio creció de 0,018 a 0,0252 meq Ca/g. La textura de las aceitunas disminuyó casi linealmente de 375 gf a 235 gf durante el tratamiento alcalino. Las aceitunas sufrieron una perdida de azúcares reductores del 25.9%. Se ha propuesto una hipotética y simplificada descripción de la dinámica de los cambios de carga iónica y el desenrrollamiento de las pectinas durante el desamarizado de las aceitunas. También se calculó el coeficiente efectivo de difusión para sodio y calcio usando un modelo de difusión para una placa compuesta. Los coeficientes resultaron en el orden de 10^{-12} m^2/s para la piel y 10^{-10} m^2/s para pulpa. En ambos casos el coeficiente de difusión de Na fue mayor que el de Ca.

PALABRAS CLAVE: Calcio – Carga iónica – Difusión – Sodio – Tratamiento alcalino de aceitunas verdes.

SUMMARY

A simple model of the diffusion phenomena taking place during the debittering process of green table olives.

The change in the concentration of sodium and calcium ions in the olive flesh and in the lye during the debittering process was quantified. The average concentration of Na increased from 0.0045 to 0,395 meq Na/g of olive flesh and the concentration of Ca increased from 0.018 to 0.0252 meq Ca/g of olive flesh. The firmness of the olives decreased almost linearly from 375 gf to 235 gf during the alkali treatment. The olives also suffered a 25.9% loss in their initial content of reducing sugars. A hypothetical simplified description of the dynamic of ionic charge changes and unwinding of the pectinic structure during the debittering process of green olives has been proposed. In addition, the effective diffusion coefficients were calculated for sodium and calcium using a diffusion model for a composite flat plate and constant diffusion coefficients. The coefficients for both solutes were in the order of 10^{-12} m^2/s for the skin and 10^{-10} m^2/s for the flesh. In both cases, the diffusion coefficients of Na were larger than the diffusion coefficients of Ca.

KEY-WORDS: Calcium – Diffusion – Debittering of green olives – Ionic charge – Sodium.

INTRODUCTION

The diffusion of solutes in food products plays an important role in many food processes, such as drying, chemical peeling, etc., as well as in the processing of fermented green olives. Given that the progress of chemical reactions that take place within the food depends on the diffusion of the reactants, these phenomena affect the progress of the processes and also the quality of the final products.

The simultaneous and individual diffusion of citric and ascorbic acid have been studied in potato tissue, cut into spheres of various diameters for different immersion times and agitation conditions. In this work the researchers determined effective diffusion coefficients for the acids diffusing individually and used multi-component analysis to evaluate interaction coefficients for the mixture of the two acids (Lombardi and Zaritzky, 1997). The effect of sodium hydroxide on the structure of pimiento peppers during peeling was evaluated using electron microscopy. The authors remarked the importance of controlling the lye concentration and exposure time in order to avoid structural damage of the tissue (Floros *et al.*, 1987).

The first step in the processing of fermented green olives is a lye treatment known as debittering. The lye treatment gives rise to complex chemical and physical changes in the olives, and its extent

also affects the subsequent diffusion of salt and the progress of the lactic fermentation (Rodríguez de la Borbolla y Alcalá, Rejano, 1979., Sciancalepore, 1984). The main objectives of this operation are to eliminate the bitter taste conferred by the glycoside oleuropein and to increase the permeability of the fruit. The skin is a natural barrier to the penetration of NaOH and other solutes to the interior of the olives. Its permeation is a function of the treatment conditions such as lye concentration and temperature, along with olive variety and maturity (Barranco *et al.*, 1987).

The effect of lye concentration (Maldonado *et al.*, 2003) and temperature (Zuritz *et al*, 2003) on the diffusion of NaOH was studied during the debittering of olives of the *Arauco variety*. The diffusion of sodium chloride into green olives placed in brines of various concentrations has also been quantified (Drusas *et al.*, 1988). In this work, the authors studied untreated olives and olives pretreated with lye at 1.8% (w/v) for 6 h and calculated salt effective diffusion coefficients assuming a hollow sphere geometry and negligible external resistance to mass transfer. They measured the absorption of salt from changes in brine concentration.

Effective diffusion coefficients of sodium during the debittering of the green olive varieties *Arauco* and *Aloreña* have also been computed (Maldonado and Zuritz, 2003, 2004). The diffusion of sodium through olive skin during and after treatment with NaOH (Zuritz and Maldonado, 2004), as well as the diffusion phenomena considering variable diffusion coefficients (Maldonado and Zuritz, 2004) have been recently studied. In both cases an increase in the skin diffusion coefficient with treatment time was observed. The same authors studied the diffusion of glucose and sodium chloride in olives as affected by different lye concentrations and different concentrations of sodium chloride (Maldonado *et al.*, 2008a, 2008b).

Texture, or fruit firmness, is significantly affected during the debittering and subsequent processing steps. The changes in texture and solubilization of cell wall polysaccharides during the processing of Spanish green olives have been extensively studied by different researchers (Jiménez *et al.*, 1995, 1996, 1997, 1998; Coimbra *et al.*, 1996). They observed a marked decrease in texture during lye treatment and washing and a noticeable increase after soaking the olives in brine. The lye treatment had the greatest effect on uronic acid-containing fractions, producing saponification of ester linkages yielding oxalate soluble polysaccharides. They indicate that electrostatic interactions and changes in the gel structure of pectic polysaccharides and the breakdown of linkages between pectic and hemicellulosic polysaccharides could be the most important factors in the loss of texture during lye treatment. The polymers with a low degree of esterification are mainly stabilized in the cell wall by ionic bonds with calcium, and therefore, an increase in their amount or ionic charge could result in a substantial recovery of the firmness

lost during the lye treatment. The softening of the olive pulp was due to a partial degradation of the pectic polysaccharides. Cell-cell adhesion is also modulated by the presence of Ca^{+2}; its removal or displacement by ions, such as Na^+, can reduce tissue firmness.

Despite the amount of studies reported on cell wall and pulp physicochemical changes during the processing of green olives, the diffusion and distribution of calcium within the pulp has not yet been quantitatively addressed, particularly in comparison with the diffusion of sodium and other solutes.

The present work presents the results obtained in the quantification of the diffusion phenomena of sodium, calcium and other components in and out of the olives during the debittering process done at industrial scale. Also, a simplified description of the dynamic of charge changes of ions is proposed to explain and complement the results being reported.

1. MATERIALS AND METHODS

1.1. Sampling

Green olives of the variety "Arauco" were used in this study. They were harvested with a Maturity Index (M.I.) 2 (Fernández *et al.*, 1991), corresponding to a green-yellow skin color, with few red spots. The M.I. scale ranges from 0 for olives with a skin color of intense green to 7 for olives with black skin and totally purple flesh. A sample of 100 olives, randomly selected from the total amount of olives chosen for processing, was measured giving the following average dimensions and standard deviations (s): weight = $5,593 \times 10^{-3}$ kg, s = 0.915×10^{-3} kg; equatorial diameter = 20.50×10^{-3} m, s = 1.59×10^{-3} m; length = 31.38×10^{-3} m, s = 2.20×10^{-3} m. A skin thickness of 4.0×10^{-5} m was obtained from a sample of 25 olives. The skin was removed with the help of a sharp thin blade, carefully eliminating any mesocarpic cells attached and avoiding cuts and breakage of the skin. The skin thickness was measured with a digital caliper Palmer Helios - 0-150 mm - Germany (1:100 mm).

The olives were placed in two large rectangular containers made of concrete, each of them holding 2100 kg of olives and 1900 liters of lye. At each sampling time; lots of twenty (20) olives and 20 ml of lye from two (2) different levels (L1: 1/3 and L2: 2/3 from the surface) were removed from each container for analysis. Each olive was pierced at the equatorial axis, all the way to the pit, with a 5-mm diameter glass tube with a sharp edge, perpendicularly to the longitudinal axis. The flesh trapped in the glass tube was then separated from the olive by cutting at the pit level. Each cylindrical sample was subsequently cut into four disks of average thickness of 1.2 mm each (measured with a 1:50 mm caliper Palmer Helios - 0-150 mm - Germany), each constituting a distinct sample for both sodium and calcium determinations.

2.2. Analytical Methods

Each disk-shaped sample was weighed in an OHAUS AP2105 (New Jersey, USA) Balance with a precision of 0.1 mg and then placed in a porcelain dish with a lid. The samples were subsequently oven dried at 100 ± 5°C for 24 h and then placed in a furnace at 550 ± 25°C until white ashes were obtained.

The ashes were quantitatively recovered with 2 ml of a 1+1 solution of HCl analytical grade purity (agp) Merck (sodium free) and 10 ml of deionized water in a test tube.

The sodium content in the above solution was measured in a flame photometer Metrolab model 315 (Buenos Aires, Argentina) by direct injection, prior to calibration of the equipment with a standard solution of NaCl (agp) Merck. The initial sodium content of the olives was determined using the same technique as mentioned above.

The content of calcium was determined with a flame atomic absorption spectrometer (FAAS) Perkin Elmer (Uberlingen, Germany) model 2380, employing the same solutions used to quantify sodium, after the addition of 5% (v/v) of a 10% (w/v) solution of strontium chloride in each sample to avoid interference from other elements. A calibration curve was prepared using a calcium standard solution made of 1.00 g/l of $Ca(NO_3)_2$ a.g.p. Tetrahedron solution and HNO_3 (agp) Merck 0,5 M. Five ml of this solution were diluted with deionized water in a 250 ml volumetric flask to achieve a calcium concentration of 25 mg/l. Thereafter, 0.25; 0.50; 0.75; 1.00; 1.50 and 2.00 ml of this solution were placed in 25 ml volumetric flasks into which 5% (v/v) of a solution of $SrCl_2$ (agp) Merck at 10% (w/v) were added and brought to volume with deionized water. A blank with deionized water and 5 % $SrCl_2$ at 10 % was used for zero adjustment.

Reducing sugars were determined in the untreated olive flesh and in lye according to Miller's technique (Maldonado et al. 2008; Miller, 1959), employing a spectrophotometer Metrolab UV-visible model 325BD (Buenos Aires, Argentina).

Changes in firmness were measured with a penetrometer Wagner (Italy); model FT01, 500 gf x 5 gf, using a 1mm diameter plunger.

2.3. Theoretical Considerations

In order to calculate the effective diffusion coefficients of both solutes into the skin (D_S) and the flesh (D_F) during the debittering process, the diffusion model for a composite flat plate and constant diffusion coefficients used by Maldonado and Zuritz (2003) was adjusted to the experimental data.

The average volumetric concentration <C> of the analytical solution to the one-dimensional diffusion process with constant effective diffusion coefficients through a composite flat plate, consisting of a thick flesh (L) and a thin skin (δ), expressed in dimensionless form, is given as (Maldonado and Zuritz, 2003):

$$\langle C \rangle = 2 \sum_{n=1}^{\infty} \frac{\sin^2(\lambda_n)}{\lambda_n \, [\lambda_n + \sin(\lambda_n) \cos(\lambda_n)]} e^{-\lambda_n^2 \theta} \qquad (1)$$

Where the λ_n are the eigen values that satisfy the following eigen function:

$$\lambda_n \tan(\lambda_n) = \frac{D_S / \delta}{D_F / L} \qquad (2)$$

In eq. (1) <C> and θ are respectively:

$$\langle C \rangle = \frac{\langle c_{(t)} \rangle - c_s}{c_i - c_s} \text{ and } \theta = \frac{D_F}{L^2} t$$

The flesh diffusivities (D_F) and eigen values λ_1 for each treatment were estimated using the least squares method, which minimizes the function:

$$S = \sum_{i=1}^{N} \left(\langle C \rangle_{exp} - \langle C \rangle_{calc} \right)^2 \qquad (3)$$

Since the sum of squares of the residuals given as equation (3) is a non-linear function of the flesh effective diffusion coefficients D_F and the eigen values λ, the iterative non-linear regression method implemented in the program Microsoft Excel Solver® was used. Next, the skin effective diffusion coefficients, D_S, were determined with equation (2).

2. RESULTS

At the beginning of the lye treatment, as the NaOH comes into contact with the skin surface, it takes some time before the lye removes the epicuticular wax and facilitates the diffusion of NaOH into the flesh. Table 1 shows the variation of the average NaOH concentration of the lye in both containers during the debittering process.

Table 1 shows that the concentration of NaOH decreased from 2.5% to a final value of 1.59% in 7.5 hours, which represents an almost 37% consumption of lye. The diffusion of the lye into

Table 1
Average lye concentration during debittering

Treatment time (hours)	Lye concentration (%)
0.0	2.50
2.5	2.07
4.5	1.82
6.5	1.64
7.5	1.59

the fruit increased the pH of the flesh from an initial value of 5 to a final value of 12. This increase in pH is an indication of the excess lye that diffused into the flesh.

The lye is an electrolytic solution where the ions sodium and hydroxyl are completely dissociated. These dissociated species have different electric charge according to their dilution. They have different degrees of solvation (Whitten and Gailey, 1988). Consequently, their affinity and capacity of combination with the olives' matrix components (pectins, fatty acids, tannins, etc) are differential and selective. They also have different particle sizes (Whitten and Gailey, 1988). Therefore, due to their selective degree of combination, reactivity and particle size, it could be assumed that their diffusivity through the olive flesh would also be different. Probably, when the hydroxyl ion diffuses into the fruit cells , it reacts forming water with hydrogen protons coming from the carboxyl groups of the fatty acids and from the hydrolysis of oleuropein, pectins, etc, partially hydrating the olive flesh. Therefore, the hydroxyl ion could be considered the true reactive ion with the components of the flesh, which disappear as such by forming water. A diffusion analysis based on the hydroxyl ion should include the corresponding rate of the reaction term in the continuity equation (Fick's Second Law of diffusion). On the other hand, even though the sodium ion also combines with fatty acids, pectins, etc., it does not transform as a species. For that reason, it can be traced inside the flesh in terms of total sodium content in each spatial dimension, regardless of whether it is in a free or combined state. In this case, a diffusion analysis based on the sodium ion does not need to include the rate of the reaction term in the continuity equation (Maldonado and Zuritz, 2003).

Based on these considerations, the following general reaction that could occur between NaOH and the fatty acids of the cuticle and the flesh cells can be proposed:
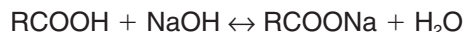
$$RCOOH + NaOH \leftrightarrow RCOONa + H_2O$$

Figure 1 shows the variation in Na^+ ion concentration within the flesh during the lye treatment. The distances represent the location of the center of each sample measured from the surface of the pit towards the skin.

The initial average concentration of Na in the fruit was 0.0045 meq Na/g of olive flesh. The figure shows a progressive increase in concentration with time in all the sections. As expected, this increase is more pronounced in the outer layer. At the end of the debittering process the average concentration of Na in the olive was 0.305 meq Na/g of olive flesh, with a minimum value of 0.200 meq Na/g of olive flesh at the inner layer in contact with the pit and of 0.395 meq Na/g of olive flesh in the outer layer in contact with the lye.

Initially, the pectins of the middle lamella in the olives would be in the form of coiled chains linked through calcium bridges (Jimenez et al., 1995). This arrangement is shown schematically in Figure 2. As indicated above, cell-cell adhesion is controlled by Ca^{+2} and its removal or displacement by Na^+ ions could reduce tissue firmness. For this reason, the content of Ca was also measured in both the pulp and the treatment liquid. The results are shown in the following figures. As mentioned before, the distances in the olive flesh represent the location
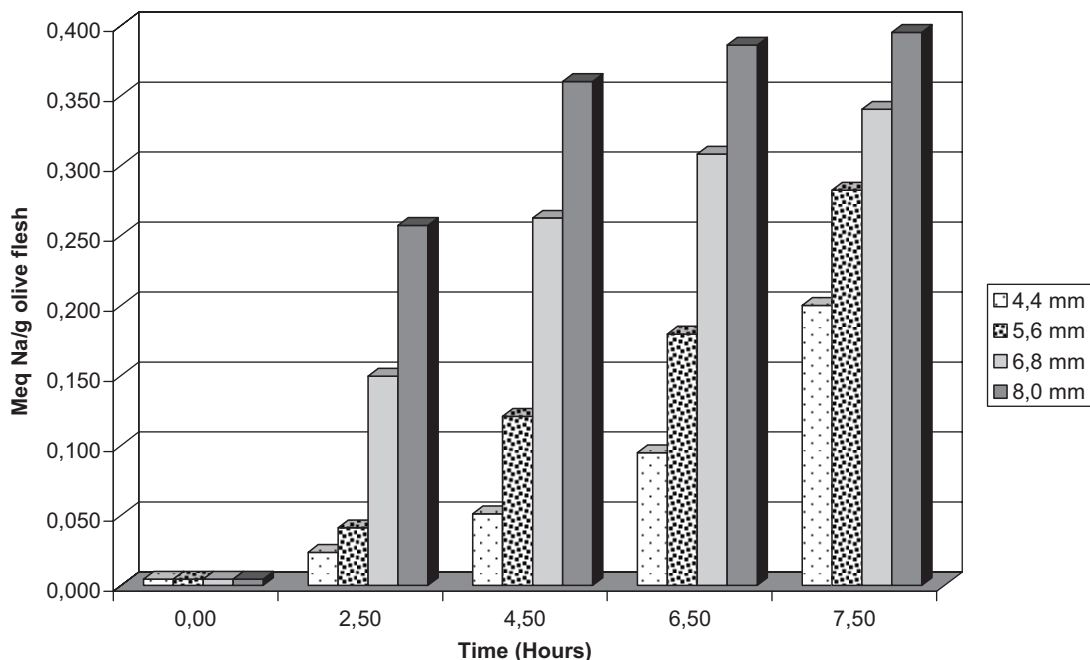

Figure 1
Concentration of sodium within the olive flesh during the debittering process. (Shown distances are from the pit outward).

of the center of each sample measured from the surface of the pit towards the skin.

Figure 3 shows the Ca concentration in the olive flesh. It can be observed that at the beginning of the debittering process, the initial average concentration of Ca was 0.018 meq Ca/g of olive flesh. Afterwards, it started to increase progressively due to the calcium concentration in the lye, following a pattern similar to that shown with Na, but with a smaller slope. The final average concentration of Ca was 0.0233 meq Ca/g of olive flesh, with a minimum value of 0.0211 meq Ca/g of olive flesh at the inner layer in contact with the pit and of 0.0252 meq Ca/g of olive flesh in the outer layer in contact with the lye.

The concentration of calcium in the lye during the debittering time is shown in Figure 4. The initial concentration of calcium in the lye was equal to 29.12 meq Ca/l. This was due to the presence of native salts such as $CaCO_3$, $Ca(CO_3H)_2$ and $CaSO_4$ in the water used to prepare the lye, typical water of the region of Mendoza (dates not shown). Thereafter, the calcium concentration continuously decreased to a final value

of 24.40 meq Ca/l. This decrease in calcium in the treatment liquid corresponds to the increase of this ion in the olive flesh.

The diffusion of Ca into the flesh was two orders of magnitude less than the diffusion of Na. For instance, while the average concentration of Ca increased with time from 0.018 to only 0.0233 meq Ca/g of olive flesh, the concentration of Na increased from 0.0045 to 0.305 meq Na/g of olive flesh. This difference in ion uptake by the olive flesh was due to the concentration gradient of each ion between the treatment liquid and the olives. The debittering solution had an initial concentration of sodium of 625 meq Na/l (NaOH at 2.5%) and of calcium equal to 29.12 meq Ca/l. Expressing these concentrations in terms of grams of olive flesh, the gradients would be respectively 0.6930 meq Na/g in the liquid versus 0.0045 meq Na/g in the olive flesh, while for calcium it would be 0.0323 meq Ca/g in the solution versus 0.0181 meq Ca/g in the olive flesh. In this context, the concentration gradient of Na was two times greater than the concentration gradient of CThe diffusion would be facilitated by the small ionic radius of these two ions, which are 0.95 A° for Na^+ and 0.99 A° for Ca^{2+} (Whitten and Gailey,1988). Nevertheless, the Ca^{2+} ion in the water can be in two different forms. On one hand, it is forming $CaCO_3$ in a colloidal state with very low solubility in water (Vogel, 1960), therefore almost not dissociated, in which case it would have to diffuse as a salt with a much larger molecular radius. And on the other hand, $Ca(CO_3H)_2$ completely dissociated as Ca^{2+} (Vogel, 1960).

Another factor to be considered is the electro negativity of the ions. At 0.93 for Na and 1.0 for Ca
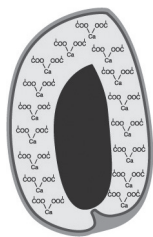

Figure 2
A simple representation of the calcium bridges of pectins of the middle lamella. Light grey represents the waxy layer. Dark grey represents the epidermis.
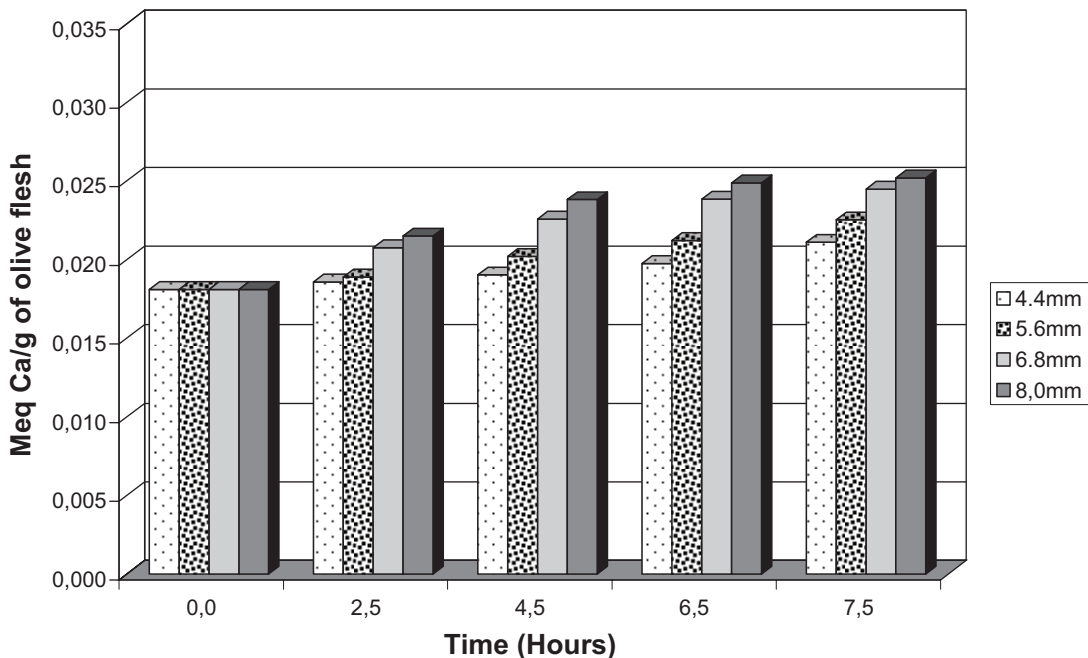

Figure 3
Concentration of calcium within the olive flesh during the debittering process. (Shown distances are from the pit outward).

(Pauling, 1932), the calcium ion would have more capacity to establish a stable linkage with COO⁻ groups from pectins than sodium ions. But the sodium ions are in larger quantity. Thus, each ion would have a different diffusion through the olive flesh and probably interact in different ways with the channel diffusion walls.

It should also be considered that the Ca⁺ ion has not only more electro negativity than the Na⁺ ion, but also a larger ionic radius that would make its diffusion into the flesh more difficult.

The effective diffusion coefficients of both ions, determined form the experimental data are shown in Table 2. For both ions, the values of the effective diffusion coefficients in the skin (D_S) are two times smaller than the values in the flesh (D_F). Furthermore, the effective diffusion coefficients for Na are slightly larger than the values for Ca, which would be in agreement with what was indicated above regarding the ionic radius of the ions and the larger quantity of sodium. These results of effective diffusion coefficients for Na are in agreement with those obtained by Maldonado and Zuritz (2003) in their laboratory scale experiments with constant lye concentrations.

As mentioned before, the initial calcium content in the olives would be responsible for the original firmness of the flesh. Figure 5 shows the evolution of firmness during the debittering process. Each value is the average of 20 readings. The firmness decreased from 375 gf to 235 gf in 7 and 1/2 hours of processing. The decrease in firmness of the olive flesh takes place in as much as the Na⁺ diffusing into the flesh reacts with the pectins and dislodges the ion Ca²⁺ (Jiménez et al., 1995, 1997). Although the olives experienced a significant softening, the concentration of calcium increased during the debittering process due to the higher Ca²⁺ concentration in the lye than in the olive flesh. Nevertheless, the Ca²⁺ dislodgment takes place due to the ionic competition between both ions as a result of the much higher Na⁺ than Ca²⁺ concentration established within the olive flesh during the debittering process.

When the olives are initially placed in the lye, the ions in the aqueous solution encounter the cuticle, a non polar lipophylic barrier, made of wax, cutine, cellulose, pectins, hydrocarbons, cratecolic acid, oleanolic acid, triterpenic alcohols, etc. (Vázquez Roncero et al., 1967), that has to be initially dissolved by the NaOH, before the diffusion process could take place in full strength. Once this step is accomplished, the diffusion flow rate would be controlled by the structure of the skin and the flesh as well as by the concentration of ions in the lye (Maldonado et al., 2003).

Table 2
Effective diffusion coefficients of Na and Ca in the olive flesh (D_F) and skin (D_S)

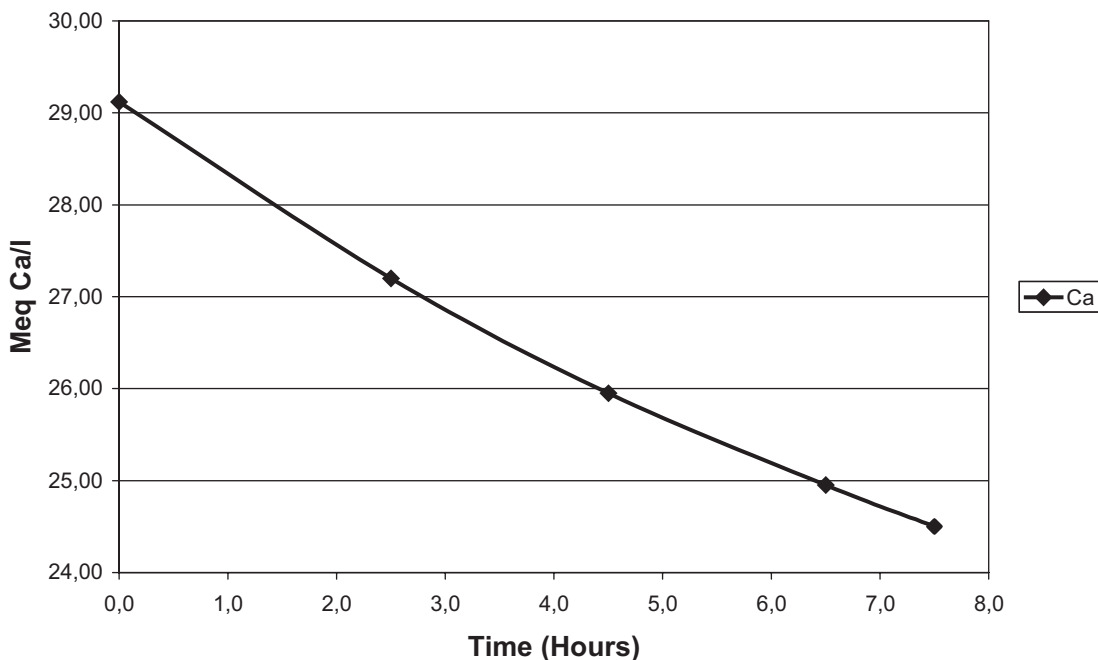| Solute	DF (m²/s)	DS (m²/s)	Regression (r²)
Sodium (Na)	6.25×10^{-10}	7.58×10^{-12}	0.956
Calcium (Ca)	5.16×10^{-10}	6.26×10^{-12}	0.994

Figure 4
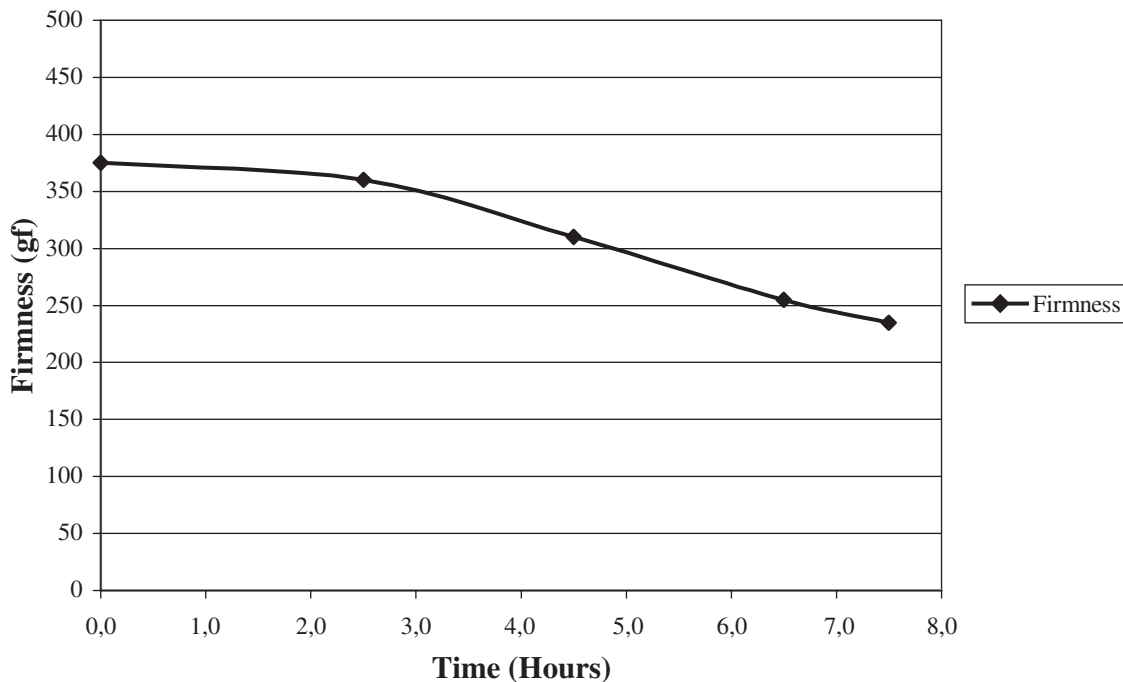Concentration of calcium in the debittering lye.

Figure 5
Evolution of firmness of olives during debittering.

As shown in Figure 6, the following simplified dynamic model of ions and charge distribution could be proposed, when the olives are initially placed in the NaOH solution. The initial pH of the olive flesh was 5, which gives an acidic medium resulting from the high concentration of fatty acids and other acids (citric, malic and oxalic acids; Fernández Díez and González Pellissó, 1956) present in the fruit. However, the positive charge of the flesh would be initially isolated from the outside by the cuticle. After the hydroxyl ions (with negative charge) have dissolved the cuticle "exposing" the flesh positive charge, the negative ions hydroxyl on the surface of the olives will react with the pectins in the cellular wall of the epidermis, forming water with the hydrogen protons of COOH groups and simultaneously opening the way to the diffusion of Na^+ ions towards the inside of the flesh and cells, where they will combine with COO^- groups of the pectins, fatty acids and other organic acids, probably establishing the charge distribution shown in Figure 7. The hydroxyl ion would be the first to react, dissolving the cuticle (shown in light grey in the Figures). Thereafter, the Na^+ ion enters into the fruit and reacts with the COO^- groups, by competition and displacing the Ca^{2+} ions that link contiguous pectic chains, causing their uncoiling. This generates a more linear and lax pectic structure and probably provokes the softening of the fruit. Although the percentage of displacement of Ca^{2+} ions is small (Jiménez et al., 1997), but apparently, it is sufficient to generate this softening, which in the present case translated to a loss of firmness from 375 gf to 235 gf. This loss of firmness associated to a disruption of the structure of the tissue would facilitate the diffusion of sugars, and other constituents such as products of the break up of the oleuropein, phenolic compounds, etc. out of the fruit.

As the hydroxyl ion reacts with the components of the cell wall, forming water with the hydrogen protons, decreasing the positive charge, there will be an increase in negative charges in the cellular structure and consequently an increase in pH to neutrality, and most of the COO^- groups would remain ionized. As was indicated previously, the excess of hydroxyl ions further increases the pH to a value of 12. This high pH will be reverted to an acidic level (pH=4.5) during the fermentation process in a subsequent step.

The change in electronic density in the flesh allows the combination of Na^+ ions to COO^- groups, generating not only pectins with a loose structure, but also the saponification of fatty acids and the break up of the oleuropein into 3,4-hydroxyfenil-ethyl alcohol or hydroxytyrosol and eleanolic acid.
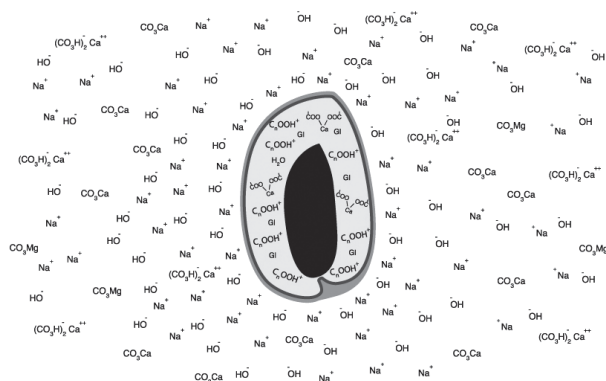

Figure 6
A simplified model of ions and charge distribution when the olives are initially placed in the
NaOH solution (Gl: glucose, CnCOOH⁺: fatty acids).

As the surface is surrounded by negative charges, the following model of charge distribution could be proposed, as shown in Figure 8.

The loss of reducing sugars during the debittering process (Fig. 9) represented an average value of 25.9% of the initial content in the olives. A fraction of the remaining 74.1% would be lost during the subsequent rinsing steps. For this reason, the debittering and the rinsing steps should be carefully controlled in order to avoid an excessive loss of sugars that could hinder the lactic fermentation process, as they are the source of carbon used by the lactic bacteria. This loss of sugars is mainly due to the structural disruption of the flesh tissue and their solubility in water.

It has been said that the alkali treatment modifies the polysaccharides of the cell wall, breaks up the weak links and probably the reducing ends are slowly degraded (Gil Martinez, 1995). In contrast with this hypothesis, we propose that what could really occur is not a degradation of the reducing ends, but rather an increase in the number of reducing molecules due to the degradation of the hydrocarbon chain in the alkali medium. In this medium, a lot of molecules of shorter length, with reducing power, can be formed from the glucose molecules. Therefore, the number of molecules with reducing power would be greater (Future studies will be undertaken in order to confirm this hypothesis).

3. CONCLUSIONS

The change in concentration of sodium and calcium ions during the debittering process was
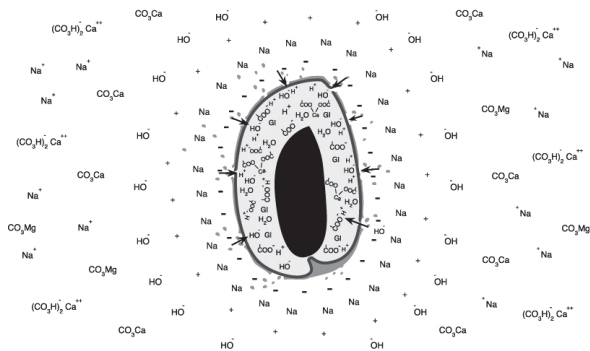

Figure 7
Position of negative and positive ions around the surface of the olives and formation of a double layer during the removal of the wax layer. The negative charges surrounding the olives represent the hydroxyl ions. Gl represents glucose. Arrows indicate entry of OH⁻ ions.
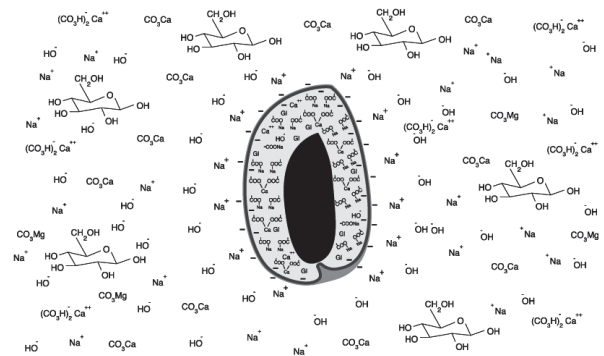

Figure 8
Simplified model of the unwinding of the pectinic structure and exit of reducing sugars into the lye during the debittering process. Gl represents glucose still remaining in the olives.
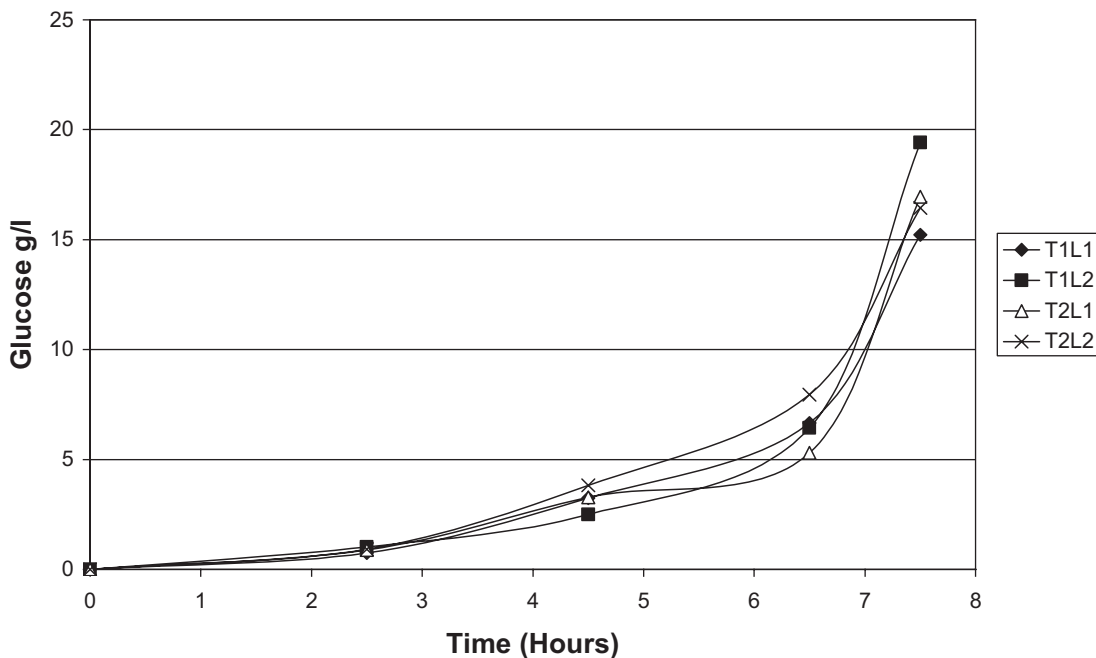

Figure 9
Loss of reducing sugars into the lye during debittering. T: represent tank and L : depth level within each tank.

quantified in the olive flesh and in the lye. Both concentrations decreased in the lye and increased in the olive flesh. The average concentration of Na increased from 0.0045 to 0.395 meq Na/g of olive flesh and the concentration of Ca increased from 0.018 to 0.0233 meq Ca/g of olive flesh. The firmness of the olives decreased almost linearly from 375 gf to 235 gf during the alkali treatment. They also suffered a 25.9% loss in reducing sugars. The values of the effective diffusion coefficients in the skin (D_S) were two times smaller than the values in the flesh (D_F), at D_S in the order of 10^{-12} and D_F in the order of 10^{-10} , respectively.

A hypothetical simplified description of the dynamic of ionic charge changes and unwinding of the pectinic structure during the debittering process of green olives has been proposed.

NOMENCLATURE

\<C\>	Dimensionless volume averaged concentration of solute.
c_i	Initial concentration of solute (meq/g olive flesh)
c_S	Solute average concentration at the outer skin surface (meq/g lye)
\<$c_{(t)}$\>	Solute concentration within the olive flesh at different time intervals (meq/g olive flesh)
D_F	Flesh effective diffusion coefficient (m^2/s)
D_S	Skin effective diffusion coefficient (m^2/s)
L	Thickness of the olive flesh (m)
t	Time (s)
δ	Skin thickness (m)
$λ_n$	Eigen values
θ	Dimensionless time

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